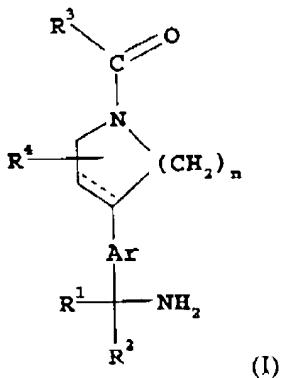
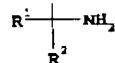


Amendments to the Claims

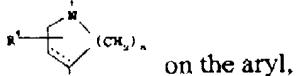
1. (Currently Amended) A compound of formula (I):-



such that Ar is an aryl group or a heteroaryl group, and the



is beta to the



wherein:-

— is a single or a double bond;

R¹ and R² are each independently hydrogen or lower alkyl;

R³ is aryl, arylalkenyl, cycloalkenyl, cycloalkyl, heteroaryl, heteroarylalkenyl, heterocycloalkenyl, a carbon linked heterocycloalkyl or alkyl optionally substituted by one or more groups selected from hydroxy, alkoxy, alkyloxycarbonylamino, cycloalkyl, heterocycloalkyl, R⁶, -OR⁶, -S(O)_mR⁶ or -C(=O)-R⁶;

R⁴ is hydrogen, acyl, alkoxy, alkyloxycarbonyl, carboxy, cyano, halo, hydroxy, -C(=O)-NY¹Y² or alkyl optionally substituted with alkoxy, alkylcarbonylamino, alkylsulfonylamino, hydroxy, -S(O)_m-alkyl or -NY¹Y²;

R⁶ is aryl or heteroaryl;

R⁷ is hydroxy, alkoxy, ureido, -C(=O)-NY¹Y², -SO₂-NY¹Y², -S(O)_m-alkyl or -NY¹Y²;

R⁸ is hydrogen or lower alkyl;

Y¹ and Y² are independently hydrogen, alkenyl, alkyl, aryl, arylalkyl, cycloalkyl, heteroaryl, heteroarylalkyl or heterocycloalkyl; or the group -NY¹Y² may form a cyclic amine;

m is zero or an integer 1 to 2; and

n is zero or an integer 1 to 4; or

an N-oxide of said compound, a prodrug of said compound, a pharmaceutically acceptable salt of said compound, a solvate of said compound, and/or a hydrate of said compound.

2. (Currently Amended) The compound of Claim 1, wherein R¹ and/or R² is hydrogen, or R¹ and R² are hydrogen, and R³ is an aryl or an heteroaryl.

3. (Currently Amended) The compound of Claim 2, wherein said R³ aryl comprises is a phenyl or a naphthyl.

4. (Currently Amended) The compound of Claim 2, wherein said R³ is aryl is substituted with at least one substituent.

5. (Original) The compound of Claim 4, wherein said substituent is selected from the group consisting of a halo atom, an alkyl substituted by aryl, an alkyl substituted by aryloxy, an alkyl substituted by aroyl, an alkyl substituted by heteroaryl, an arylalkynyl, a heteroarylalkynyl, an aryl, a heteroaryl, an arylalkenyl and an arylalkyloxy.

6. (Original) The compound of Claim 5, wherein said aryl or heteroaryl of said substituent is further substituted by at least one aryl group substituent.

7. (Currently Amended) The compound of Claim 2, wherein said heteroaryl comprises is a pyridyl, a quinolinyl, a thienyl, a furanyl, or an indolyl.

8. (Original) The compound of Claim 7, wherein said heteroaryl is substituted with at least one substituent.

9. (Currently Amended) The compound of Claim 8, wherein said substituent comprises is an alkyl, an alkyl substituted by an aryl, an alkyl substituted by an aryloxy, an alkyl substituted by an aroyl, an alkyl substituted heteroaryl, an arylalkynyl, a heteroarylalkynyl, a heteroaryl, an arylalkenyl or an arylalkyloxy.

10. (Original) The compound of Claim 9, wherein said aryl of said substituent is further substituted by at least one aryl substituent.

11. (Currently Amended) The compound of Claim 1, wherein R⁴ comprises is hydrogen or a cyano group.

12. (Currently Amended) The compound of Claim 1, wherein R⁵ comprises is a hydrogen, a lower alkyl, or a halo.

13. (Cancelled) The compound of Claim 1, wherein _____ is a single bond.

14. (Original) The compound of Claim 1, wherein n=2.

15. (Currently Amended) The compound of Claim 1, wherein:

Ar comprises is a phenyl group;

R¹=R²=hydrogen;

R¹ and R² are both hydrogen;

R³ comprises is an aryl, a naphthyl or a heteroaryl;

R⁴ comprises is hydrogen or a cyano;

_____ is a single bond; and

_____ is a single bond; and

n=2.

16. (Currently Amended) The compound of Claim 15, wherein R³ said as aryl or said naphthyl of R³-is substituted with at least one substituent comprising selected from the group consisting of a halo atom, an alkyl substituted by aryl, an alkyl substituted by aryloxy, an alkyl substituted by aroyl, an alkyl substituted by aryloxy, an alkyl substituted by aroyl, an alkyl substituted by a heteroaryl, an arylalkynyl, a heteroarylkynyl, an aryl, a heteroaryl, an arylalkenyl, or and an arylalkyloxy.

17. (Original) The compound of Claim 16, wherein said aryl or said heteroaryl of said substituent is further substituted by at least one aryl substituent.

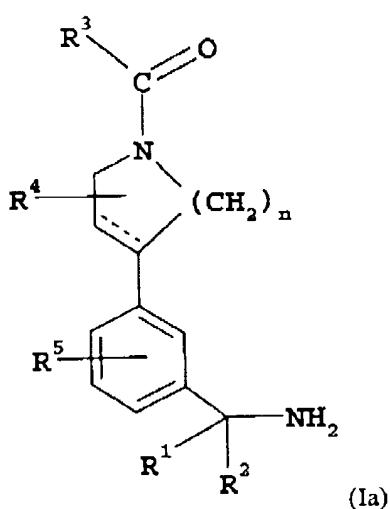
18. (Currently Amended) The compound of Claim 15, wherein R³ said as heteroaryl of R³ is substituted by at least one substituent comprising selected from the group consisting of a pyridyl, a quinolinyl, a thienyl a furanyl, or and an indolyl.

19. (Currently Amended) The compound of Claim 18, wherein said substituent of said heteroaryl is further substituted by at least one moiety comprising selected from the group

consisting of an alkyl substituted by an aryl, an alkyl substituted by an aryloxy, an alkyl substituted by an aroyl, an alkyl substituted heteroaryl, an arylalkynyl, a heteroarylalkynyl, a heteroaryl, an arylalkenyl, or and an arylalkyloxy.

20. (Original) The compound of Claim 19, wherein an aryl of said moiety is further substituted by at least one aryl substituent.

21. (Currently Amended) A compound of formula (Ia):



wherein

R¹ and R² are each independently hydrogen or lower alkyl;

R³ is aryl, arylalkenyl, cycloalkenyl, cycloalkyl, heteroaryl, heteroarylalkenyl, heterocycloalkenyl, a carbon linked heterocycloalkyl or alkyl optionally substituted by one or more groups selected from hydroxy, alkoxy, alkyloxycarbonylamino, cycloalkyl, heterocycloalkyl, R⁶, -OR⁶, -S(O)_mR⁶ or -C(=O)-R⁶;

R⁴ is hydrogen, acyl, alkoxy, alkyloxycarbonyl, carboxy, cyano, halo, hydroxy, -C(=O)-NY¹Y² or alkyl optionally substituted with alkoxy, alkylcarbonylamino, alkylsulfonylamino, hydroxy, -S(O)_m-alkyl or -NY¹Y²; and

R⁵ is hydrogen, acyl, alkoxy, alkyloxycarbonyl, aryl, carboxy, cyano, halo, heteroaryl, heteroaryloxy, heterocycloalkyl, heterocycloalkyloxy, heterocycloalkylalkyloxy, heteroarylalkyloxy, hydroxy, trifluoromethyl, -C(=O)-NY¹Y², -NY¹Y²,

-Z¹-C₂₋₆alkylene-R⁷ or alkyl optionally substituted with alkoxy, alkylcarbonylamino, alkylsulfonylamino, aryl, heteroaryl, heterocycloalkyl, hydroxy, ureido, -C(=O)-NY¹Y², -SO₂-NY¹Y², -S(O)_m-alkyl or -NY¹Y², and, and

R⁶ is aryl or heteroaryl;

R⁷ is hydroxy, alkoxy, ureido, -C(=O)-NY¹Y², -SO₂-NY¹Y², -S(O)_m-alkyl or -NY¹Y²,

R⁸ is hydrogen or lower alkyl;

Y¹ and Y² are independently hydrogen, alkenyl, alkyl, aryl, arylalkyl, cycloalkyl, heteroaryl, heteroarylalkyl or heterocycloalkyl; or the group -NY¹Y² may form a cyclic amine;

Z¹ is O, S(O)_m or NR⁸;

m is zero or an integer 1 to 2; and

n is zero or an integer 1 to 4; or

a corresponding an N-oxide of said compound, a prodrug of said compound, a pharmaceutically acceptable salt of said compound, a solvate of said compound, an N-oxide of said solvate of said compound, or a prodrug of said solvate of said compound N-oxides and prodrugs.

22. (Currently Amended) The compound of Claim 21, wherein R³ is an aryl comprising a phenyl or a naphthyl.

23. (Currently Amended) The compound of Claim 22, wherein said aryl is substituted by at least one substituent comprising selected from the group consisting of a halo atom, an alkyl substituted by an aryl, and an alkyl substituted by a heteroaryl.

24. (Original) The compound of Claim 23, wherein said aryl or heteroaryl of said substituent is further substituted by at least one aryl group substituent.

25. (Currently Amended) The compound of Claim 21, wherein R³ comprises is phenylC₁₋₃alkylpyridyl [e.g. 5-phenylethyl-pyrid-3-yl], phenylC₁₋₃alkylthienyl [e.g. 5-phenylethyl-thien-2-yl] or indolyl [e.g. indol-6-yl].

26. (Currently Amended) The compound of Claim 21, wherein R³ is a heteroaryl comprising selected from the group consisting of a pyridyl, a quinolinyl, a thienyl, a furanyl, or and an indolyl.

27. (Currently Amended) The compound of Claim 26, wherein said heteroaryl is substituted by at least one substituent comprising selected from the group consisting of an alkyl substituted by an aryl, or and an alkyl substituted by a heteroaryl.

28. (Original) The compound of Claim 27, wherein said aryl and said heteroaryl of said substituent are further substituted by at least one aryl group substituent.

29. (Currently Amended) The compound of Claim 28, wherein R³ is phenylC₁-alkylpyridyl [e.g. 5-phenylethyl pyrid-3-yl], phenylC₁-alkylthienyl [e.g. 5-phenylethyl-thien-2-yl] or indolyl [e.g. indol-6-yl].

30. (Currently Amended) The compound of Claim 21, wherein R⁴ comprises is a hydrogen or a cyano.

31. (Currently Amended) The compound of Claim 21, wherein R⁵ comprises is a hydrogen, a lower alkyl or a halo.

32. (Currently Amended) The compound of Claim 31, wherein R⁵ comprises is methyl or fluoro.

33. (Original) The compound of Claim 31, wherein R⁵ is attached to the phenyl ring of formula (Ia) in the position para to the CH₂NH₂ group.

34. (Currently Amended) The compound of Claim 21, wherein:
R³ is a phenyl, a naphthyl, a heteroaryl selected from the group consisting a pyridyl, a quinolinyl, a thienyl, a furanyl, and an indolyl, a phenyl substituted by at least one substituent, a naphthyl substituted by at least one substituent, or a heteroaryl heteroaryl selected from the group consisting a pyridyl, a quinolinyl, a thienyl, a furanyl, and an indolyl, that is substituted by at least one substituent,

wherein said substituent is selected from the group consisting of a halo atom, an alkyl substituted by aryl, and alkyl substituted heteroaryl, wherein the aryl or heteroaryl groups are further substituted by one or more aryl group substituents;

R⁴ comprises is hydrogen or a cyano; and

R⁵ comprises is hydrogen, a lower alkyl or a halo.

35. (Currently Amended) The compound of Claim 34, wherein:

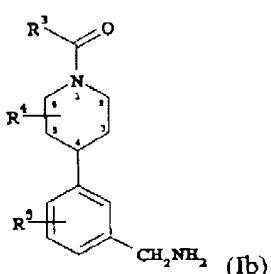
R^3 comprises is phenylC₁₋₃alkylpyridyl [e.g. 5-phenylethyl pyrid-3-yl], phenylC₁₋₃alkylthienyl [e.g. 5-phenylethyl thien-2-yl] or indolyl [e.g. indol-6-yl];

R^4 comprises is a hydrogen or a cyano; and

R^5 comprises is a methyl or a fluoro, and is attached to the phenyl ring of formula (Ib) (la) in the position para to the $CR^1R^2NH_2$, CH_2NH_2 group.

36. (Currently Amended) The compound of Claim 21, having A compound of formula

(Ib):



wherein

R^3 is aryl, arylalkenyl, cycloalkenyl, cycloalkyl, heteroaryl, heteroarylalkenyl, heterocycloalkenyl, a carbon-linked heterocycloalkyl or alkyl optionally substituted by one or more groups selected from hydroxy, alkoxy, alkylcarboxylamino, cycloalkyl, heterocycloalkyl, R^6 , OR^6 , $S(O)_mR^6$ or $C(=O)R^6$;

R^4 is hydrogen, acyl, alkoxy, alkylcarboxyl, carboxy, cyano, halo, hydroxy, $-C(=O)NY^1Y^2$ or alkyl optionally substituted with alkoxy, alkylcarboxylamino, alkylsulfonylamino, hydroxy, $S(O)_m$ alkyl or NY^1Y^2 ; and

R^5 is hydrogen, acyl, alkoxy, alkylcarboxyl, aryl, carboxy, cyano, halo, heteroaryl, heteroaryloxy, heterocycloalkyl, heterocycloalkyloxy, heterocycloalkylalkyloxy, heteroarylalkyloxy, hydroxy, trifluoromethyl, $-C(=O)NY^1Y^2$, NY^1Y^2 , and

$-Z^1-C_2-alkylene-R^7$ or alkyl optionally substituted with alkoxy, alkylcarboxylamino, alkylsulfonylamino, aryl, heteroaryl, heterocycloalkyl, hydroxy, ureido, $C(=O)NY^1Y^2$, $-SO_2-NY^1Y^2$, $-S(O)_m$ alkyl or NY^1Y^2 ; and,

a corresponding N-oxide of said compound, a prodrug of said compound, a pharmaceutically acceptable salt of said compound, a solvate of said compound, an N-oxides and prodrugs.

37. (Currently amended) The compound of Claim 36, wherein R³ as is an aryl comprising is selected from the group consisting of a phenyl or and a naphthyl.

38. (Currently amended) The compound of Claim 37, wherein said aryl is substituted by at least one substituent comprising selected from the group consisting of a halo atom, an alkyl substituted by an aryl, and an alkyl substituted by a heteroaryl.

39. (Original) The compound of Claim 38, wherein said aryl or heteroaryl of said substituent is further substituted by at least one aryl group substituent.

40. (Currently amended) The compound of Claim 36, wherein R³ is a heteroaryl comprising pyridyl, a quinolinyl, a thienyl, a furanyl, or an indolyl.

41. (Currently Amended) The compound of Claim 41, wherein said heteroaryl is substituted by at least one substituent comprising selected from the group consisting of an alkyl substituted by an aryl, or and an alkyl substituted by a heteroaryl.

42. (Original) The compound of Claim 42, wherein said aryl and said heteroaryl of said substituent are further substituted by at least one aryl group substituent.

43. (Currently amended) The compound of Claim 36, wherein R³ comprises is phenylC₁₋₃alkylpyridyl [e.g. 5-phenylethyl-pyrid-3-yl], phenylC₁₋₃alkylthienyl [e.g. 5-phenylethyl-thien-2-yl] or indolyl [e.g. indol-6-yl].

44. (Currently amended) The compound of Claim 36, wherein R⁴ comprises is a hydrogen or a cyano.

45. (Currently amended) The compound of Claim 36, wherein R⁵ comprises is a hydrogen, a lower alkyl or a halo.

46. (Currently amended) The compound of Claim 45, wherein R⁵ comprises is a methyl or a fluoro.

47. (Original) The compound of Claim 45, wherein R⁵ is attached to the phenyl ring of formula (Ib) in the position para to the CH₂NH₂ group.

48. (Currently amended) The compound of Claim 36, wherein:

R³ is a phenyl, a naphthyl, a heteroaryl selected from the group consisting a pyridyl, a quinolinyl, a thienyl, a furanyl, and an indolyl, a phenyl substituted by at least one substituent, a naphthyl substituted by at least one substituent, or a heteroaryl heteroaryl selected from the group consisting a pyridyl, a quinolinyl, a thienyl, a furanyl, and an indolyl, that is substituted by at least one substituent,

wherein said substituent is selected from the group consisting of a halo atom, an alkyl substituted by aryl, and alkyl substituted heteroaryl, wherein the aryl or heteroaryl groups are further substituted by one or more aryl group substituents;

R⁴ comprises is hydrogen or a cyano; and

R⁵ comprises is hydrogen, a lower alkyl or a halo.

49. (Currently amended) The compound of Claim 48, wherein:

R³ is phenylC₁₋₃alkylpyridyl [e.g. 5-phenylethyl-pyrid-3-yl], phenylC₁₋₃alkylthienyl [e.g. 5-phenylethyl-thien-2-yl] or indolyl [e.g. indol-6-yl];

R⁴ comprises is a hydrogen or a cyano; and

R⁵ comprises is a methyl or a fluoro.

50. (Original) The compound of Claim 1, selected from the group consisting of:

3-[1-(5-phenylethynyl-pyridine-3-carbonyl)-piperidin-4-yl]-benzylamine;

3-[1-(3-phenylethyl-benzoyl)-piperidin-4-yl]-benzylamine;

3-{1-[3-(4-hydroxyphenyl)ethyl-benzoyl]-piperidin-4-yl}-benzylamine;

3-{1-[3-(6-amino-pyridin-3-yl)ethyl-benzoyl]-piperidin-4-yl}-benzylamine;

3-[1-(5-phenylethyl-thiophene-2-carbonyl)-piperidin-4-yl]-benzylamine;

4-fluoro-3-[1-(5-phenylethyl-pyridine-3-carbonyl)-piperidin-4-yl]-benzylamine;

4-methyl-3-[1-(5-phenylethyl-pyridine-3-carbonyl)-piperidin-4-yl]-benzylamine;

3-[1-(indole-6-carbonyl)-piperidin-4-yl]-benzylamine;

4-(3-aminomethyl-phenyl)-1-(5-phenethyl-pyridine-3-carbonyl)-piperidine-4-carbonitrile

[4-(3-aminomethylphenyl)piperidin-1-yl]-[3,4-dichlorophenyl]methanone;

1-[1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl]-3-methylsulfanyl-6,7-dihydro-

5H-benzo[c]thiophen-4-one trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(3-methylsulfanyl-6,7-dihydro-benzo[c]thiophen-1-yl)-methanone trifluoroacetate;

1-{1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl}-3-ethylsulfanyl-6,6-dimethyl-6,7-dihydro-5H-benzo[c]thiophen-4-one trifluoroacetate;

1-{1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl}-3-propylsulfanyl-6,7-dihydro-5H-benzo[c]thiophen-4-one trifluoroacetate;

1-{1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl}-3-isopropylsulfanyl-6,7-dihydro-5H-benzo[c]thiophen-4-one trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-benzo[b]thiophen-2-yl-methanone-trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-4-hydroxy-piperidin-1-yl]-1-(5-phenethyl-pyridin-3-yl)-methanone-ditrifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(1-methyl-1H-indol-3-yl)-methanone-trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-[3-(2-fluoro-phenylethynyl)-phenyl]-methanone trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-{3-[2-(2-fluoro-phenyl)-ethyl]-phenyl}-methanone trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-{3-[2-(6-amino-pyridin-3-yl)-ethyl]-phenyl}-methanone tri-trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(6-chloro-thieno[3,2-*b*]thiophen-2-yl)-methanone trifluoroacetate;

(3R,4S) and (3S, 4R)-4-(3-Aminomethyl-phenyl)-1-(5-phenethyl-pyridine-3-carbonyl)-piperidine-3-carboxylic acid ethyl ester dihydrochloride;

3-[1-(5-Phenylethynyl-furan-2-carbonyl)-piperidin-4-yl]-benzylamine trifluoroacetate;

4-(3-Aminomethyl-phenyl)-piperidine-1-carboxylic acid (3,4-dichloro-phenyl)-amide trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(2,3-dihydro-benzofuran-5-yl)-methanone;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(5,6-dichloro-pyridin-3-yl)-methanone;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(3-bromo-4-fluoro-phenyl)-methanone;

(E)-1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-3-(2-nitro-phenyl)-propenone;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(3-bromo-5-iodo-phenyl)-methanone; and

(E)-1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-3-phenyl-propenone.

51. (Currently Amended) A pharmaceutical composition comprising a pharmaceutically acceptable amount of a compound of Claim 1 and a pharmaceutically acceptable carrier thereof.

52. (Original) The pharmaceutical composition of Claim 51, wherein said compound is selected from the group consisting of:

3-[1-(5-phenylethynyl-pyridine-3-carbonyl)-piperidin-4-yl]-benzylamine;
3-[1-(3-phenylethyl-benzoyl)-piperidin-4-yl]-benzylamine;
3-{1-[3-(4-hydroxyphenyl)ethyl-benzoyl]-piperidin-4-yl}-benzylamine;
3-{1-[3-(6-amino-pyridin-3-yl)ethyl-benzoyl]-piperidin-4-yl}-benzylamine;
3-[1-(5-phenylethyl-thiophene-2-carbonyl)-piperidin-4-yl]-benzylamine;
4-fluoro-3-[1-(5-phenylethyl-pyridine-3-carbonyl)-piperidin-4-yl]-benzylamine;
4-methyl-3-[1-(5-phenylethyl-pyridine-3-carbonyl)-piperidin-4-yl]-benzylamine;
3-[1-(indole-6-carbonyl)-piperidin-4-yl]-benzylamine;
4-(3-aminomethyl-phenyl)-1-(5-phenethyl-pyridine-3-carbonyl)-piperidine-4-carbonitrile
[4-(3-aminomethylphenyl)piperidin-1-yl]-(3,4-dichlorophenyl)methanone;
1-{1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl}-3-methylsulfanyl-6,7-dihydro-
5H-benzo[c]thiophen-4-one trifluoroacetate;
1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(3-methylsulfanyl-6,7-dihydro-
benzo[c]thiophen-1-yl)-methanone trifluoroacetate;
1-{1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl}-3-ethylsulfanyl-6,6-dimethyl-
6,7-dihydro-5H-benzo[c]thiophen-4-one trifluoroacetate;
1-{1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl}-3-propylsulfanyl-6,7-dihydro-
5H-benzo[c]thiophen-4-one trifluoroacetate;
1-{1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-methanoyl}-3-isopropylsulfanyl-6,7-
dihydro-5H-benzo[c]thiophen-4-one trifluoroacetate;
1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-benzo[b]thiophen-2-yl-methanone-
trifluoroacetate;
1-[4-(3-Aminomethyl-phenyl)-4-hydroxy-piperidin-1-yl]-1-(5-phenethyl-pyridin-3-yl)-
methanone-ditrifluoroacetate;
1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(1-methyl-1H-indol-3-yl)-methanone-
trifluoroacetate;
1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-[3-(2-fluoro-phenylethynyl)-phenyl]-
methanone trifluoroacetate;

1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-{3-[2-(2-fluoro-phenyl)-ethyl]-phenyl}-methanone trifluoroacetate;
1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-{3-[2-(6-amino-pyridin-3-yl)-ethyl]-phenyl}-methanone tri-trifluoroacetate;
1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(6-chloro-thieno[3,2-*b*]thiophen-2-yl)-methanone trifluoroacetate;
(3R,4S) and (3S, 4R)-4-(3-Aminomethyl-phenyl)-1-(5-phenethyl-pyridine-3-carbonyl)-piperidine-3-carboxylic acid ethyl ester dihydrochloride;
3-[1-(5-Phenylethynyl-furan-2-carbonyl)-piperidin-4-yl]-benzylamine trifluoroacetate;
4-(3-Aminomethyl-phenyl)-piperidine-1-carboxylic acid (3,4-dichloro-phenyl)-amide trifluoroacetate;
1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(2,3-dihydro-benzofuran-5-yl)-methanone;
1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(5,6-dichloro-pyridin-3-yl)-methanone;
1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(3-bromo-4-fluoro-phenyl)-methanone;
(E)-1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-3-(2-nitro-phenyl)-propenone;
1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-1-(3-bromo-5-iodo-phenyl)-methanone;
(E)-1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-3-phenyl-propenone; and
1-[4-(3-Aminomethyl-phenyl)-piperidin-1-yl]-3-cyclohexyl-propan-1-one.

53. (Currently Amended) A pharmaceutical composition comprising a pharmaceutically acceptable amount of a compound of Claim 21 and a pharmaceutically acceptable carrier thereof.

54. (Currently Amended) A pharmaceutical composition comprising a pharmaceutically acceptable amount of a compound of Claim 36 and a pharmaceutically acceptable carrier thereof.

55. (Currently Amended) A method for treating asthma in a patient ~~suffering from, or subject to, a condition that can be ameliorated by the administration of an inhibitor of trypsin, wherein the method comprises administering to the patient an a pharmaceutically effective amount of a compound of Claim 1.~~

56. (Cancelled)

57. (Cancelled)

58. (Currently Amended) A method for treating asthma in a patient suffering from, or subject to, a condition that can be ameliorated by the administration of an inhibitor of trypsin, wherein the method comprises administering to the patient an a pharmaceutically effective amount of a compound of Claim 21.

59. (Cancelled)

60. (Cancelled)

61. (Currently Amended) A method for treating asthma in a patient suffering from, or subject to, a condition that can be ameliorated by the administration of an inhibitor of trypsin, wherein the method comprises administering to the patient an a pharmaceutically effective amount of a compound of Claim 36.

62. (Cancelled)

63. (Cancelled)

64. (Withdrawn) A pharmaceutical composition comprising a compound of Claim 1 and a second compound selected from the group consisting of a beta adrenergic agonist, an anticholinergic, an anti-inflammatory corticosteroid, and an anti-inflammatory agent; and a pharmaceutically acceptable carrier thereof.

65. (Withdrawn) The pharmaceutical composition of Claim 64, wherein the beta adrenergic agonist comprises albuterol, terbutaline, formoterol, fenoterol or prenaline; the anticholinergic comprises ipratropium bromide; the anti-inflammatory corticosteroid comprises beclomethasone dipropionate, triamcinolone acetonide, flunisolide or dexamethasone; and the anti-inflammatory agent comprises sodium cromoglycate or nedocromil sodium.

66. (Withdrawn) A pharmaceutical composition comprising a compound of formula 21 and a second compound selected from the group consisting of a beta adrenergic agonist, an anticholinergic, an anti-inflammatory corticosteroid, and an anti-inflammatory agent; and a pharmaceutically acceptable carrier thereof.

67. (Withdrawn) The pharmaceutical composition of Claim 66, wherein the beta andrenergic agonist comprises albuterol, terbutaline, formoterol, fenoterol or prenalin; the anticholinergic comprises ipratropium bromide; the anti-inflammatory corticosteroid comprises beclomethasone dipropionate, triamcinolone acetonide, flunisolide or dexamethasone; and the anti-inflammatory agent comprises sodium cromoglycate or nedocromil sodium.

68. (Withdrawn) A pharmaceutical composition comprising a compound Claim 36 and a second compound selected from the group consisting of a beta andrenergic agonist, an anticholinergic, an anti-inflammatory corticosteroid, and an anti-inflammatory agent; and a pharmaceutically acceptable carrier thereof.

69. (Withdrawn) The pharmaceutical composition of Claim 66, wherein the beta andrenergic agonist comprises albuterol, terbutaline, formoterol, fenoterol or prenalin; the anticholinergic comprises ipratropium bromide; the anti-inflammatory corticosteroid comprises beclomethasone dipropionate, triamcinolone acetonide, flunisolide or dexamethasone; and the anti-inflammatory agent comprises sodium cromoglycate or nedocromil sodium.

70. (Withdrawn) A method for treating a patient suffering from asthma, comprising administering to the patient a combination of a compound of Claim 1, and a second compound selected from the group consisting of a beta andrenergic agonist, an anticholinergic, an anti-inflammatory corticosteroid, and an anti-inflammatory agent

71. (Withdrawn) A method for treating a patient suffering from asthma, comprising administering to the patient a combination of a compound of Claim 21, and a second compound selected from the group consisting of a beta andrenergic agonist, an anticholinergic, an anti-inflammatory corticosteroid, and an anti-inflammatory agent.

72. (Withdrawn) A method for treating a patient suffering from asthma, comprising administering to the patient a combination of a compound of Claim 36, and a second compound selected from the group consisting of a beta andrenergic agonist, an anticholinergic, an anti-inflammatory corticosteroid, and an anti-inflammatory agent.

73. (New) The compound of Claim 25, wherein the phenylC₁₋₃alkylpyridyl is 5-phenylethyl-pyrid-3-yl, the phenylC₁₋₃alkylthienyl is and the indolyl is indol-6-yl.

74. (New) The compound of Claim 29, wherein the phenylC₁₋₃alkylpyridyl is 5-phenylethyl-pyrid-3-yl, the phenylC₁₋₃alkylthienyl is 5-phenylethyl-thien-2-yl and the indolyl is indol-6-yl.

75. (New) The compound of Claim 35, wherein the phenylC₁₋₃alkylpyridyl is 5-phenylethyl-pyrid-3-yl, the phenylC₁₋₃alkylthienyl is 5-phenylethyl-thien-2-yl and the indolyl is indol-6-yl.

76. (New) The compound of Claim 1, wherein ~~-----~~ is a single bond.